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         Sep 29
NEWS
      2
                 The Philippines Inventory of Chemicals and Chemical
                 Substances (PICCS) has been added to CHEMLIST
NEWS
         Oct 27
                 New Extraction Code PAX now available in Derwent
                 Files
NEWS
         Oct 27
                 SET ABBREVIATIONS and SET PLURALS extended in
                 Derwent World Patents Index files
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         Oct 27
                 Patent Assignee Code Dictionary now available
                 in Derwent Patent Files
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         Oct 27
                 Plasdoc Key Serials Dictionary and Echoing added to
                 Derwent Subscriber Files WPIDS and WPIX
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      7
         Nov 29
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         Dec 5
                 French Multi-Disciplinary Database PASCAL Now on STN
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         Dec 15
                 2001 STN Pricing
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         Dec 17
                 Merged CEABA-VTB for chemical engineering and
                 biotechnology
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                 Corrosion Abstracts on STN
NEWS 13
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FULL ESTIMATED COST
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0.15

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=> lipoprotein

L1 354253 LIPOPROTEIN

=> lysing agent

L2 221 LYSING AGENT

=> luekocytes

L3 55 LUEKOCYTES

=> 11 and 12

L4 0 L1 AND L2

=> high density lipoprotein

L5 83404 HIGH DENSITY LIPOPROTEIN

=> 11 and 13

L6 0 L1 AND L3

=> fixing white blood cells

7 FILES SEARCHED...

L7 0 FIXING WHITE BLOOD CELLS

=> fixing cells

L8 459 FIXING CELLS

=> 18 and blood cell?

2 FILES SEARCHED... 3 FILES SEARCHED...

L9 11 L8 AND BLOOD CELL?

=> dupo rem 19

MISSING OPERATOR REM L9
The search profile that was entered contains terms or nested terms that are not separated by a logical operator.

=> dup rem 19

PROCESSING COMPLETED FOR L9
L10 4 DUP REM L9 (7 DUPLICATES REMOVED)

=> d 1-4 ibib ab

L10 ANSWER 1 OF 4 MEDLINE

ACCESSION NUMBER: 1999451073 MEDLINE

DOCUMENT NUMBER: 99451073

TITLE: Key adhesion molecules are present on long podia extended

DUPLICATE 1

by hematopoietic cells.

AUTHOR: Holloway W; Martinez A R; Oh D J; Francis K; Ramakrishna

R;

Palsson B O

CORPORATE SOURCE: Department of Bioengineering, University of California at

San Diego, La Jolla, California.

CONTRACT NUMBER: R01 HL59234 (NHLBI)

RO1 HL60398 (NHLBI)

SOURCE: CYTOMETRY, (1999 Nov 1) 37 (3) 171-7.

Journal code: D92. ISSN: 0196-4763.

PUB. COUNTRY: United States

Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 200001 ENTRY WEEK: 20000104

AB BACKGROUND: We recently reported that CD34(+) hematopoietic cells and the KGla cell line extend long, thin podia. These podia can dynamically

extend

and retract, often adhere to the substrate, and appear to connect cells

up

to 300 &mgr;m apart. The surface receptors found on these podia have not been described. METHODS:By using time-lapse fluorescent microscoscopy and immunostaining techniques, we describe a method for detecting surface receptors on these podia. This includes an in situ antibody staining procedure without fixing cells. RESULTS:We demonstrate, using CD34 selected mobilized peripheral blood cells and KGla cells, that adhesion molecules known to play important roles in blood-cell migration and adhesion are present on these podia. These include: CD11a, CD18, CD29, CD34, CD45, CD49d, CD49e, and CD62L. Additionally, CD54 and CD44 were present on the podia extended by KGla cells, but were not detectable on the primary CD34(+) cells. The integrin CD49d localized at the base of these podia in a time-dependent manner in KGla cells. The frequency and morphology of these long podia on three myeloid leukemia-cell lines (KG1a, MV4-11, and AML-193) and a CD34-negative T-cell line (CEM) are also compared. KG1a

and

CEM cell lines extend long, dynamic podia that are similar to the podia

primary CD34(+) cells in morphology and adhesion molecule expression. The AML-193 and MV4-11 cell lines, however, did not extend these long podia. CONCLUSIONS: We describe a technique that provides a method of detecting surface receptors on thin cell membrane projections. These results support

the likely role of these podia in cell migration and cell-cell communication. Copyright 1999 Wiley-Liss, Inc.

L10 ANSWER 2 OF 4 EMBASE COPYRIGHT 2001 ELSEVIER SCI. B.V.

ACCESSION NUMBER: 1998147335 EMBASE

TITLE:

Reactivity of workshop monoclonal antibodies on

paraformaldehyde-fixed porcine blood mononuclear cells.

AUTHOR:

Schuberth H.-J.; Rabe H.-U.; Leibold W.

CORPORATE SOURCE:

H.-J. Schuberth, Immunology Unit, School of Veterinary

Medicine, Bischofsholer Datum 15, D-30173 Hannover,

Germany. jschub@immunologie.tiho-hannover.de

SOURCE:

Veterinary Immunology and Immunopathology, (30 Jan 1998)

60/3-4 (409-417).

Refs: 15

ISSN: 0165-2427 CODEN: VIIMDS

PUBLISHER IDENT.:

S 0165-2427(97)00115-3

COUNTRY:

Netherlands

DOCUMENT TYPE:

Journal; Conference Article

FILE SEGMENT:

General Pathology and Pathological Anatomy 005

026 Immunology, Serology and Transplantation

LANGUAGE:

English

SUMMARY LANGUAGE:

English

One hundred sixty-four monoclonal antibodies (mAbs) of the second international swine CD workshop were tested for their reactivity with porcine blood mononuclear cells before and after fining the cells with varying concentrations of paraformaldehyde (PFA) (1, 5 and 10 g l-1). A total of 38 (out of 134) positive reacting mAbs were significantly affected in their binding behavior on fixed cells. Modulation was seen as reduction in binding (staining intensity and/or % positive cells, n = 18) or in elevated values (n = 20). Modified mAb binding occurred after fixing cells with 5 to 10 g 1-1 PFA.

L10 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2001 ACS

DUPLICATE 2

ACCESSION NUMBER:

1993:511109 CAPLUS

DOCUMENT NUMBER:

119:111109

TITLE:

Process for analyzing clastogenic agents

INVENTOR(S):

Tometsko, Andrew M.

PATENT ASSIGNEE(S):

Litron Laboratories, USA

SOURCE:

U.S., 29 pp.

DOCUMENT TYPE:

CODEN: USXXAM

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

KIND

PATENT INFORMATION:

PATENT NO.

DATE APPLICATION NO.

US 1990-492584

19900313

Α 19930720 Potential clastogenic agent is identified by analyzing the change in AB micronucleated cells. The invention discloses the procedures for dosing mice, obtaining blood samples, fixing and staining cells, configuring the flow cytometer condition for micronuclei anal., the mode of data acquisition, and anal. The malarial parasite, Plasmodium berghei, provides an excellent model for optimizing cell fixing, cell staining,

and

instrument calibration. The cells are fixed at ultralow temps. to provide

cells suitable for staining and high speed flow cytometry anal. Good results are obtained when Hoechst 33258 is used as a DNA stain and propidium iodide is used as the RNA stain. Assays were done using methylmethane sulfonate and cyclophosphamide as clastogenic agents in

L10 ANSWER 4 OF 4 MEDLINE

DUPLICATE 3

ACCESSION NUMBER:

76136251

MEDLINE

DOCUMENT NUMBER:

76136251

TITLE:

The scanning electron microscopy of normal human

peripheral

blood lymphocytes.

AUTHOR:

Newell D G; Roath S; Smith J L

SOURCE:

BRITISH JOURNAL OF HAEMATOLOGY, (1976 Mar) 32 (3) 309-16.

Journal code: AXC. ISSN: 0007-1048.

PUB. COUNTRY:

ENGLAND: United Kingdom

Journal; Article; (JOURNAL ARTICLE)

LANGUAGE:

English

FILE SEGMENT:

Priority Journals

ENTRY MONTH:

197607

A study of the effects of various sample preparation techniques for scanning electron microscopy has been undertaken in an attempt to resolve conflicting descriptions of the surface topography of human peripheral blood lymphocytes. By fixing cells in suspension--a

technique thought most likely to avoid the production of artefacts--no clearly defined morphological classification of lymphocytes could be

made.

and when T- and B-lymphocyte enriched preparations were studied their surfaces appeared similar. Both T- and B-rosetted cells showed identical morphological changes as a result of their interaction with red blood cells. The smooth cells described in other reports were found only under certain conditions of preparation. It is therefore not possible to distinguish between T- and B-cell populations, using the S.E.M., on the basis of surface morphology alone.

=> d his

(FILE 'HOME' ENTERED AT 16:12:27 ON 23 MAR 2001)

FILE 'MEDLINE, BIOSIS, EMBASE, CEABA-VTB, CABA, LCA, CAPLUS, CA' ENTERED AT 16:12:37 ON 23 MAR 2001

L1354253 LIPOPROTEIN 221 LYSING AGENT

L2 L3 55 LUEKOCYTES

0 L1 AND L2

83404 HIGH DENSITY LIPOPROTEIN

0 L1 AND L3

L7 O FIXING WHITE BLOOD CELLS

rs459 FIXING CELLS

L9 11 L8 AND BLOOD CELL?

L10 4 DUP REM L9 (7 DUPLICATES REMOVED)

=> composition?

2 FILES SEARCHED...

3922722 COMPOSITION?

=> 111 and cell? and fixing 3 FILES SEARCHED... 1226 L11 AND CELL? AND FIXING => 112 and lys? agent 0 L12 AND LYS? AGENT => 112 and lytic agent 0 L12 AND LYTIC AGENT => 112 and lipoprotein? 2 L12 AND LIPOPROTEIN? => d 1-2 ibib ab L15 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2001 ACS ACCESSION NUMBER: 1993:621153 CAPLUS DOCUMENT NUMBER: 119:221153 TITLE: ' Hematology control composition for leukocyte analogs and methods for their preparation and use INVENTOR(S): Young, Carole; Elliott, Michael N.; Fischer, Timothy J.; Naylor, Nancy R. PATENT ASSIGNEE(S): Coulter Corp., USA PCT Int. Appl., 49 pp. CODEN: PIXXD2 DOCUMENT TYPE: Patent

SOURCE:

LANGUAGE: English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

	PATENT NO.				KIND DATE				APPLICATION NO.				DATE			
	WO	9317 W:			A: CA,	1 1993 JP, KR,	30902 NO,		WO	1993-	 -US185	- - 5	1993	0217		
		RW:	AT,	BE,	CH,	DE, DK,	ES,	FR,	GB,	GR, IE	I. IT.	LU.	MC.	NI.	PT.	SE
	ΑU	9337	366		A.	1 1993	30913			1993-					,	
	EP	6281	67		A.	1994	1214			1993-			19930			
	EΡ		.67			l 2000								_		
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	US	5320	964		Α		0614		US	1993-	81752		19930	0623		
	NO	9403	115		À	1994	0901		NO	1994-	3115		19940	0823		
PRIO	RITY	APP	LN.	INFO.	:				US	1992-	84043		1992			
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mprising leukocyte analogs is described. The

analogs comprise red blood cells which simulate .gtoreq.2 phys. properties of human leukocytes. A method for making leukocyte analogs from blood cells having desired phys. properties is also described. The process comprises expanding the cell vol., changing the Hb content of the cell, and fixing the cell. Generally, the monocyte and lymphocyte analogs leak Hb from

the cell while the eosinophil analog has the Hb pptd. in the cell. A further method is described to use the control product to det. whether an automatic instrument is operating within the manufacturer's specification. Lymphocyte analogs were prepd. from goose red blood cells and monocyte, eosinophil, and neutrophil analogs were prepd. from alligator red blood cells. The analogs were resuspended in an aq. soln. of Moducyte contg. cholesterol. This assembly

could be stored for up to .apprx.6 mo with the addn. of known stabilizers.

L15 ANSWER 2 OF 2 CA COPYRIGHT 2001 ACS ACCESSION NUMBER: 119:221153 CA

TITLE:

INVENTOR(S):

Hematology control composition for leukocyte analogs and methods for their preparation and use

Young, Carole; Elliott, Michael N.; Fischer, Timothy

J.; Naylor, Nancy R.

PATENT ASSIGNEE(S): SOURCE:

Coulter Corp., USA PCT Int. Appl., 49 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

	PA	TENT NO.		KIN	ND DATE		APPLICATION NO. DATE
					19930902		WO 1993-US1855 19930217
		W: AU RW: AI			JP, KR, NO, DE, DK, ES,		GB, GR, IE, IT, LU, MC, NL, PT, SE
		9337366	5	A1	19930913		AU 1993-37366 19930217
		628167		A1	19941214		EP 1993-906274 19930217
	EΡ	628167		В1	20000823		
		R: AT	BE,	CH,	DE, DK, ES,	FR,	GB, GR, IE, IT, LI, LU, MC, NL, PT,
SE							•
		0750403			19950427		JP 1993-515120 19930217
	BR	9305952		Α	19971021		BR 1993-5952 19930217
	AT	195810		E	20000915		AT 1993-906274 19930217
	US	5320964		Α	19940614		US 1993-81752 19930623
	NO	9403115		Α	19940901		NO 1994-3115 19940823
PRIO	RITY	APPLN.	INFO	. :			US 1992-840435 19920224
							WO 1993-US1855 19930217
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A hematol. control product comprising leukocyte analogs is described. AΒ The

analogs comprise red blood cells which simulate .gtoreq.2 phys. properties of human leukocytes. A method for making leukocyte analogs from blood cells having desired phys. properties is also described. The process comprises expanding the cell vol., changing the Hb content of the cell, and fixing the cell. Generally, the monocyte and lymphocyte analogs leak Hb from the cell while the eosinophil analog has the Hb pptd. in the cell. A further method is described to use the control product to det. whether an automatic instrument is operating within the manufacturer's specification. Lymphocyte analogs were prepd. from goose red blood cells and monocyte, eosinophil, and neutrophil analogs were prepd. from alligator red blood cells. The analogs were resuspended in an aq. soln. of Moducyte contg. cholesterol. This

could be stored for up to .apprx.6 mo with the addn. of known stabilizers.

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